Patent claims

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- A vector for inserting a nucleic acid into a cell, which vector contains a low molecular weight polyethylenimine (LMW PEI) and a nucleic acid, with the LMW PEI having a molecular weight of less than 50,000 Da.
 - 2. A vector as claimed in claim 1, wherein the LMW PEI has a molecular weight of from 500 to 30,000 Da.

3. A vector as claimed in either of claim 1 and 2, wherein the LMW PEI has a molecular weight of from 1000 to 5000 Da.

4. A vector as claimed in one or more of claims 1 to 3, wherein the LMW PEI has a molecular weight of about 2000 Da.

- 5. A vector as claimed in one or more of claims 1 to 4, wherein the nucleic acid is a viral or nonviral nucleic acid construct.
- 20 6. A vector as claimed in ene or more of claims 1 te-5, wherein the nucleic acid construct contains one or more effector genes.
 - 7. A vector as claimed in one or more of claims 1 to 6, wherein at least one effector gene encodes a pharmacological active compound or its prodrug form.

8. A vector as claimed in one or more of claims 1 to 7, wherein at least one effector gene encodes an enzyme.

9. A vector as claimed in one-or-more-of-claims 1 to-8, wherein at least one effector gene is expressed together with a cell-specific ligand as a fusion protein.

Claim 1

10. A vector as claimed in one or more of claims 1 to 9, wherein the LMW PEI is coupled to a cell-specific ligand.

11. A vector as claimed in one or more of claims 1 to 10, wherein the cell-specific ligand binds to the outer membrane of a target cell.

A vector as claimed in one or more of claims 12. target cell is an endothelial cell, a muscle cell, a macrophage, a lymphocyte, a glia cell, an hematopoietic cell, a tumor cell, a virusinfected cell, a bronchial epithelial cell or a liver cell. 5 A vector as claimed in one or more of claims 1 to 12, wherein the 13.

ratio by weight of LMW PEI to nucleic acid is 3:1 or more.

A vector as claimed in one or more of claims 1 to 13, wherein the 14. 10 ratio by weight of LMW PEI to nucleic acid is 8:1 or more.

> 15. A process for preparing a low molecular weight polyethylenimine (LMW PEI) having a molecular weight of less than 50,000 Da. which comprises monomeric ethylenimine being polymerized in aqueous solution by adding hydrochloric acid.

> The process as claimed in claim 15, wherein the aqueous solution is 16. from 0.1% strength to 90% strength with respect to monomeric ethylenimine and from 0.1% strength to 10% strength with respect to concentrated hydrochloric acid.

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Claim 15
The process as claimed in either of claims 15 and 16; wherein the 17. polymerization is carried out at a reaction temperature of from 30°C to 70°C.

The process as claimed in one or more of claims 15 to 17; wherein 18. the reaction time is from 1 to 30 days.

19. A low molecular weight polyethylenimine which has a molecular weight of less than 50,000 Da and which is prepared by a process according to one or more of claims 15 to 18.

20. The use of a low molecular weight polyethylenimine having a molecular weight of less than 50,000 Da for preparing a vector as claimed in one or more of claims 1 to 14:

Claim I 21. A process for preparing a vector according to one or more of claims. -1 to 14, which comprises mixing an appropriate quantity of

LMW PEI with an appropriate quantity of nucleic acid in an aqueous

			solution.
J	5	22.	The use of a vector as claimed in one or more of claims 1 to 14 for inserting a nucleic acid into a cell.
	10	23.	The use of a vector as claimed in claim 22, wherein the cell is an endothelial cell, a lymphocyte, a macrophage, a liver cell, a fibroblast, a muscle cell or an epithelial cell.
	10	24.	A process for preparing a transfected cell, which comprises incubating a vector as claimed in one or more of claims 1 to 14 in vitro with this cell.
	15	25.	Claim 1 A transfected cell which contains a vector as claimed in one or more of claims 1 to 14.
	20	26.	The use of a transfected cell as claimed in claim 25 for preparing a pharmaceutical.
		27.	The use of a low molecular weight polyethylenimine as claimed in claim 19 for preparing a pharmaceutical.
	25	28.	The use of a vector as claimed in ene or more of claims 1 to 14-for preparing a pharmaceutical.
2	٠	29.	The use of a vector as claimed in one or more of claims 1 to 14-for preparing a pharmaceutical for gene therapy.
	30	30.	A process for preparing a pharmaceutical, which comprises mixing a nucleic acid with an LMW PEI.
C		31.	A pharmaceutical which comprises a vector as claimed in ene or more of claims 1 to 14.
	35	32.	A pharmaceutical which comprises an LMW PEI as claimed in claim 19.

33. A pharmaceutical which comprises a transfected cell as claimed in claim 25.